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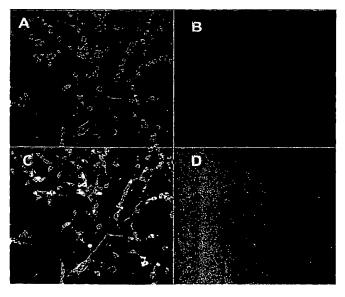
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(54) Title: MONOCLONAL ANTIBODY IMAGING AND THERAPY OF TUMORS THAT EXPRESS MET AND BIND HEP-ATOCYTE GROWTH FACTOR



(57) Abstract: In a wide variety of human solid tumors, an aggressive, metastatic phenotype and poor clinical prognosis are associated with expression of the receptor tyrosine kinase Met and its agonist ligand HGF. Disclosed herein are (a) mAbs and hybridoma cell lines that produce them, which mAbs antibodies are specific for Met and (b) combinations of anti-Met and anti-HGF mAbs. When detectably labeled, these antibodies are useful for imaging such tumors. Anti-Met mAb compositions and methods for scintigraphic detection, diagnosis, prognosis, monitoring and therapy of Met-bearing tumors are provided.



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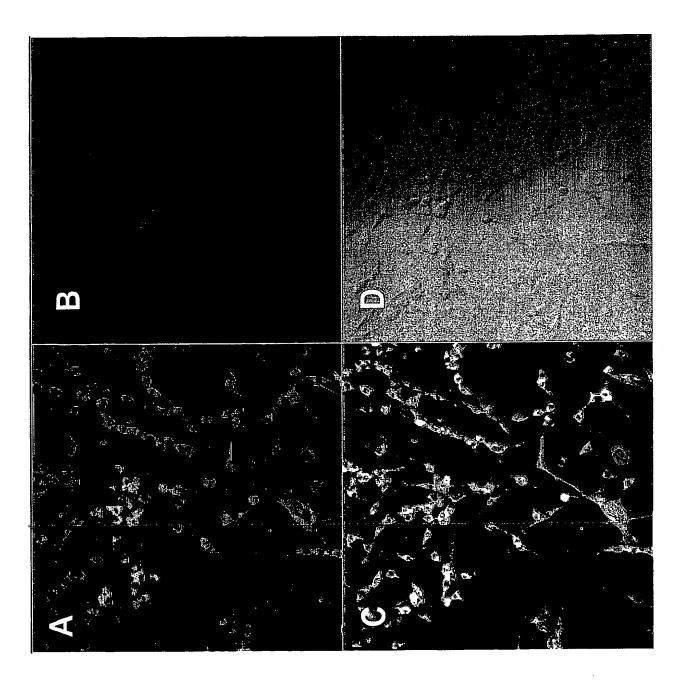
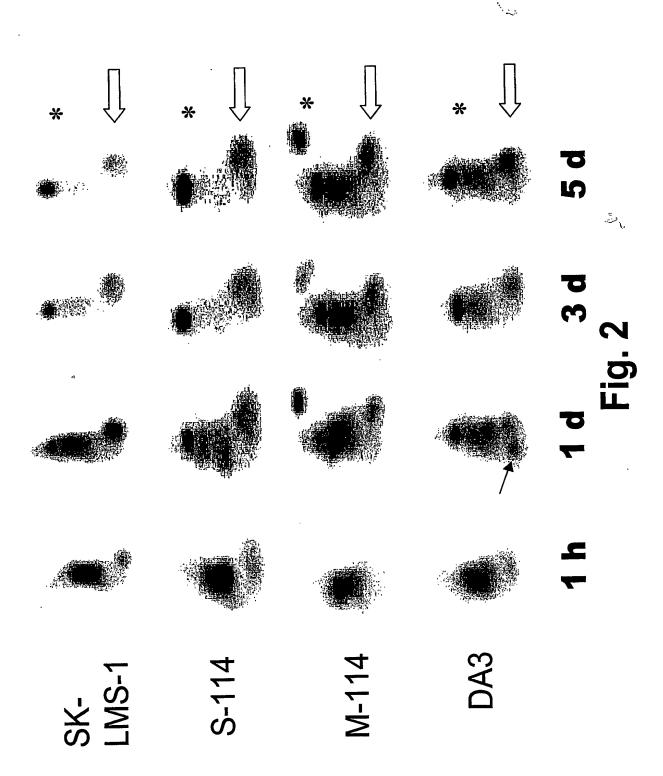
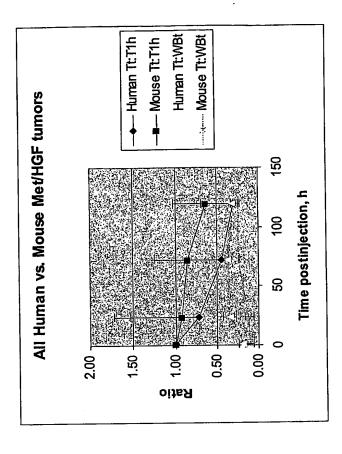


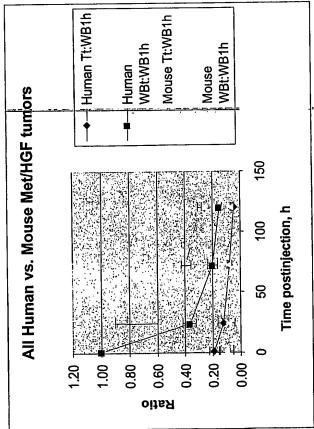
Fig. 1



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# Potential mechanisms of \*MAb binding to tumor cells

Direct cell targeting with \*anti-Met

Indirect cell targeting with \*anti-HGF/SF

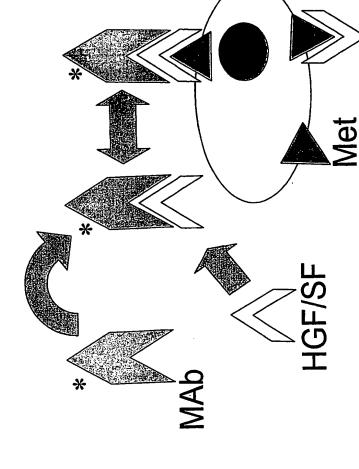
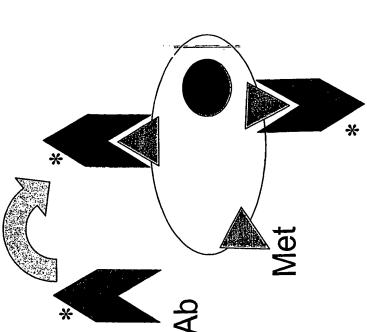
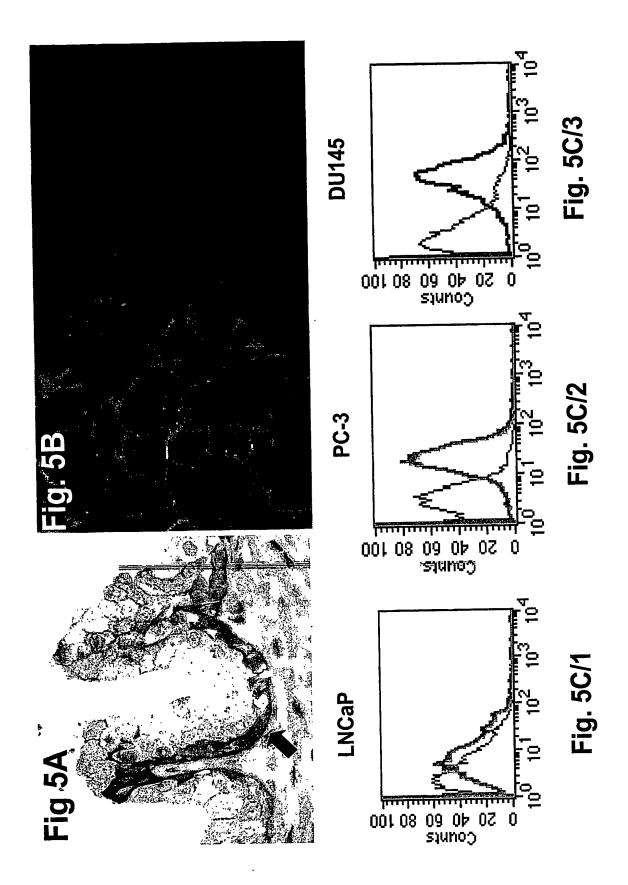
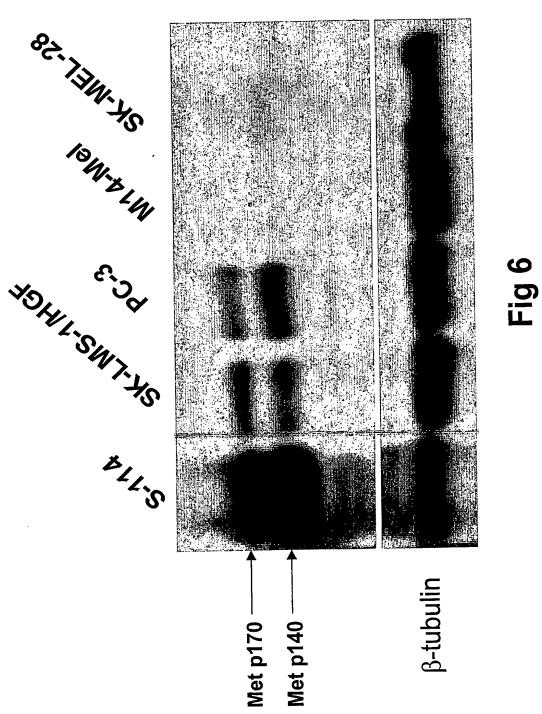
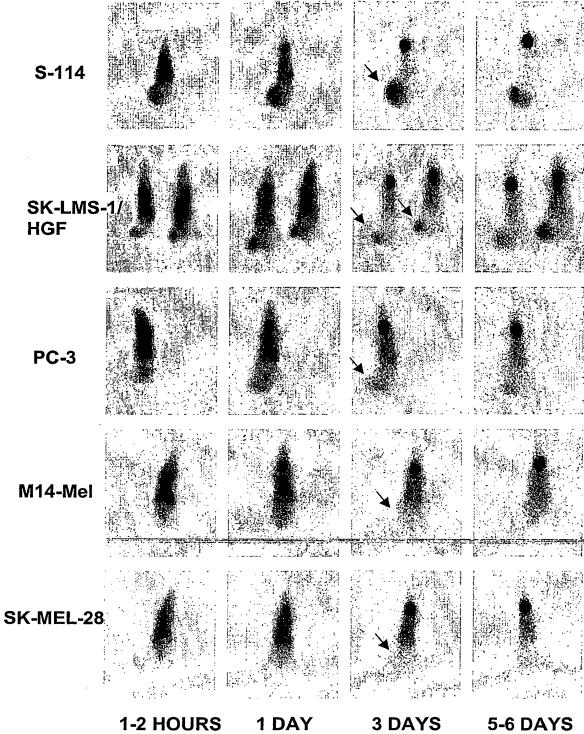


Fig. 4

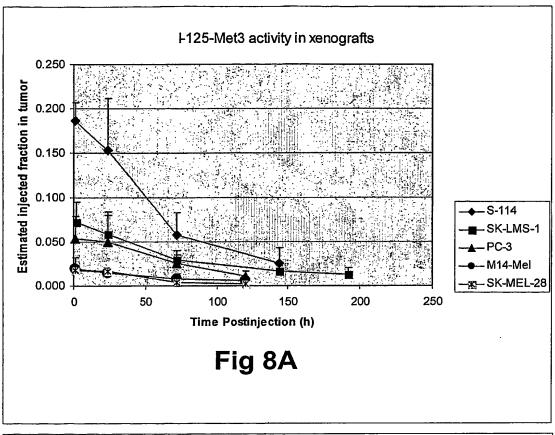


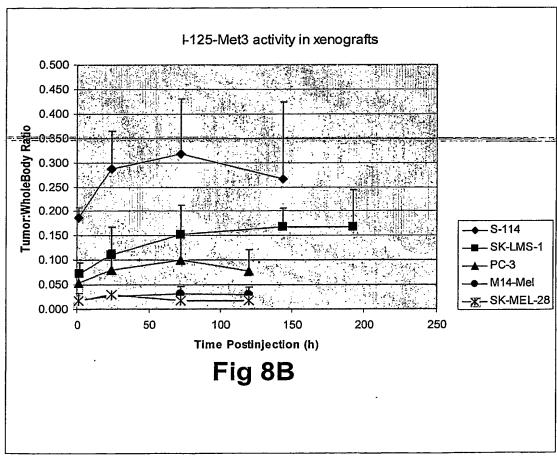




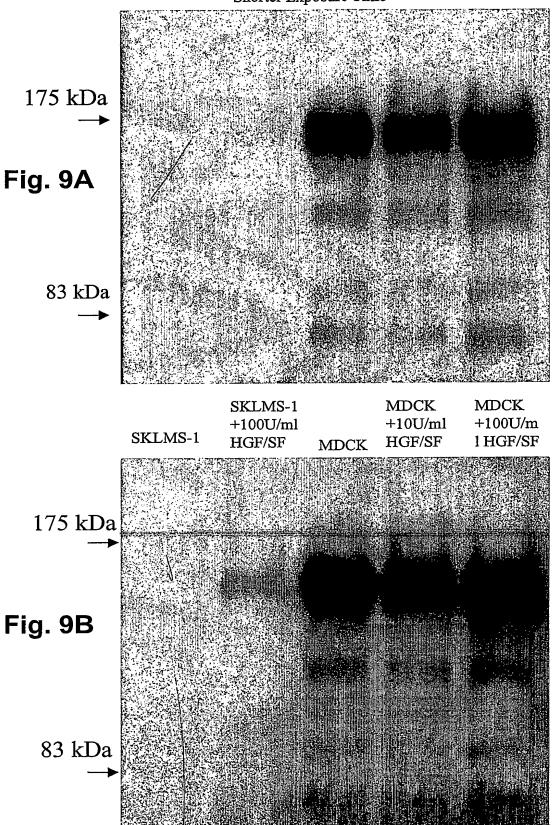


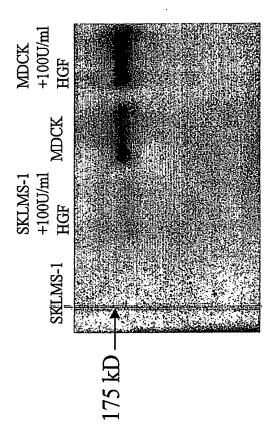
1-2 HOURS 1 DAY 3 DAYS 5-6 DAYS
Time post-injection
Fig. 7





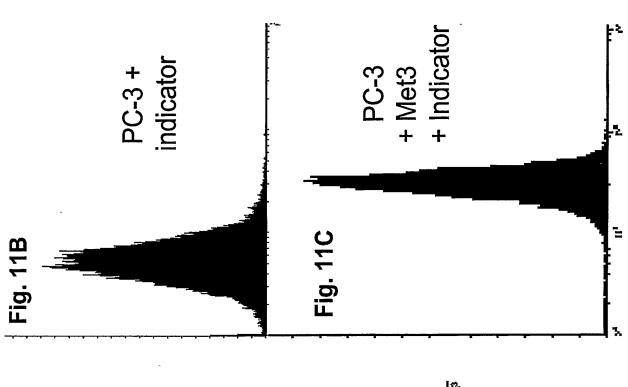
# Shorter Exposure Time

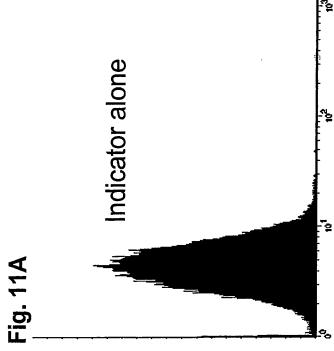


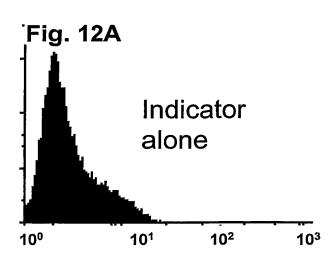


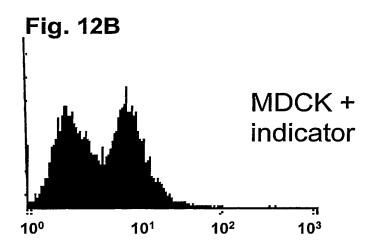
Met5 immunoprecipitates

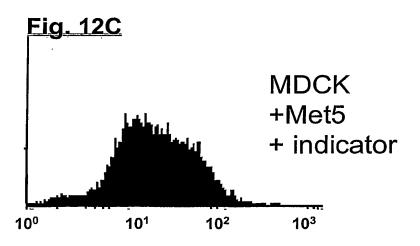
Fig. 10

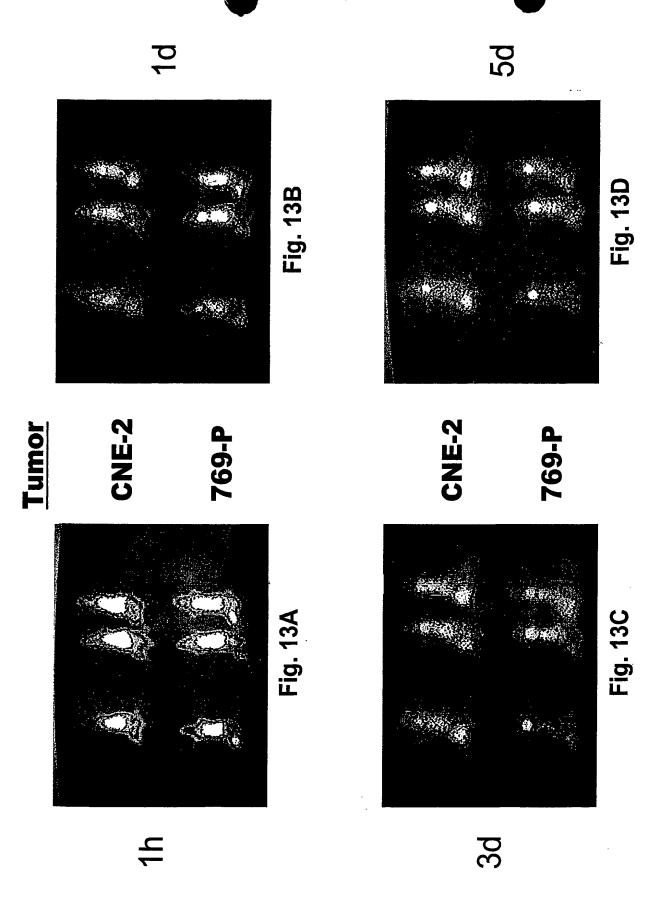












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